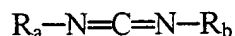


CLAIMS

1. (Currently Amended) A method for preparing a steroidal carbothiolic acid or a salt thereof, said method comprises:

A) reacting a steroidal carboxylic acid or a salt thereof with a coupling agent selected from the group consisting of carbodiimide derivatives represented by the following formula:



wherein  $R_a$  and  $R_b$  are the same or different, and each represent an aliphatic, heteroaliphatic, carbocyclic or a heterocyclic group, wherein the group is ~~all said groups are~~ optionally substituted};

alone or in conjunction with a coupling enhancer; and

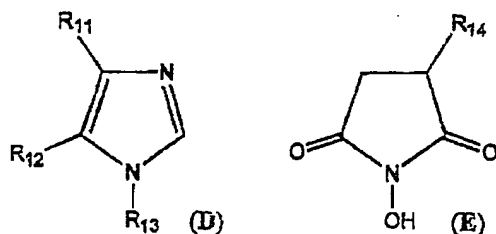
B) reacting the product of step A) with a nucleophilic agent comprising a sulfur atom.

2. (Original) A method according to claim 1 in which the coupling agent is 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide (EDC).

3. (Original) A method according to claim 2, in which the coupling agent is the hydrochloride salt of EDC.

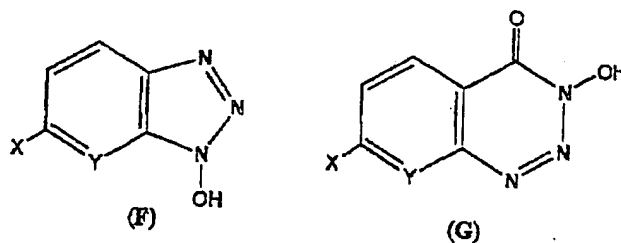
4. (Currently Amended) A method according to claim 1 any of the preceding claims, in which the coupling enhancer is selected from the group consisting of:

A) a heterocyclic ring containing one or two nitrogen atoms, said ring being optionally substituted; such as a compound of formula (D) or formula (E),



wherein  $R_{11}$  and  $R_{12}$  can be the same or different, and each represent a hydrogen atom or a cyano group;  $R_{13}$  represent a hydrogen atom or an alkyl group; and  $R_{14}$  represent a hydrogen atom or a salt of a sulfonic acid such as sodium sulfonate [ $S(=O)(=O)O^-Na^+$ ]; and

B) an unsaturated 5-6 membered heterocyclic ring fused to an aromatic or heteroaromatic ring in which the said heterocyclic ring contains three nitrogen atoms, said rings being optionally substituted, such as a compound of formula (F) or formula (G),



$X = H, F, Cl, Br$  and  $Y = CH, N, O, S$

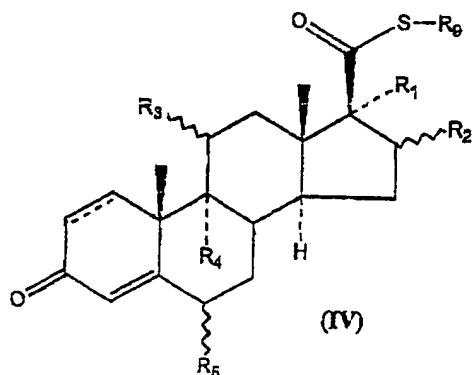
Preferably 6-chloro-hydroxybenzotriazole (6-Cl-HOBT), 7-aza-hydroxybenzotriazole (HOAt), or 3-hydroxy-4-ene-3,4-dihydro-1,2,3-benzotriazine (Dbht-OH).

5. (Currently Amended) A method according to claim 1 ~~any of the preceding claims~~, where the nucleophilic agent comprising a sulfur atom is selected from the group consisting of ~~comprising~~:

compounds of formula  $[M]^+ [SH]^-$  wherein M is a metal such as Li, Na or K; or  $[M]^{2+} [S]^{2-}$  wherein M is a metal such as Ca or Mg, the said sulfide salts being optionally hydrated ~~(such as sodium hydrosulfide hydrate)~~; and  
an *in situ* generated sulfide salt or a hydrated sulfide salt.

6. (Currently Amended) The method of claim 1 ~~any of the preceding claims~~, wherein the nucleophilic agent is dissolved in a suitable solvent prior to addition to the reaction mixture, or wherein the nucleophilic agent is added in the form of a solid salt or as a solution of the salt in water, ~~and/or~~ an organic solvent, or a combination thereof.

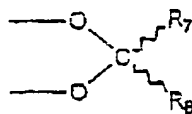
7. (Currently Amended) A method according to claim 1 ~~any of the preceding claims~~ for preparing a steroidal carbothioic acid of formula (IV) or a salt thereof



wherein the symbol  $\equiv$  in the 1,2-position represent a single or a carbon-carbon double bond;

$R_1$  represents a hydrogen atom, a hydroxy- or an alkoxy group (such as an optionally substituted  $C_{1-6}$ -alkoxy) in the  $\alpha$ -configuration, a group  $-O-C(=O)-R_6$  is an alkyl group (such as optionally substituted  $C_{1-5}$ -alkyl) or an optionally substituted 5-6 membered heterocyclic ring containing either oxygen, nitrogen or sulfur as ring hetero atom (such as a furanyl, pyrrolyl or a thiophenyl group);

$R_2$  represents a hydrogen atom, a hydroxy group, an alkoxy group (such as an optionally substituted  $C_{1-6}$ -alkoxy) in the  $n$ -configuration, an alkyl group (such as an optionally substituted  $C_{1-6}$ -alkyl) which may be in either the  $\eta$ - or  $\beta$ -configuration, an alkylene group (such as an optionally substituted  $C_{1-6}$ -alkylene having the two free valencies on the same carbon atom preferably methylene), wherein [the alkylene group is bound to the steroid nucleus via a double bond,] or  $R_1$  and  $R_2$  together represent



where  $R_7$  and  $R_8$  are the same or different and each represent a hydrogen atom or an alkyl group (such as an optionally substituted  $C_{1-6}$  alkyl);

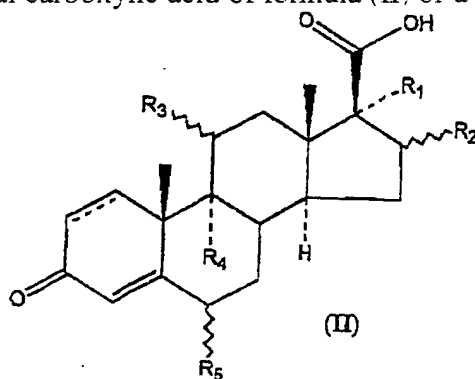
$R_3$  represent a hydrogen atom, hydroxy- or a protected hydroxy group in either a  $\alpha$ - or  $\beta$ -configuration or an oxo group (in which case the bond between  $R_3$  and the steroid nucleus is a double bond);

$R_4$  represents a hydrogen- or a halogen atom or  $R_3$  and  $R_4$  together represent a carbon-carbon bond or an epoxy group in the  $\beta$ -configuration; and

$R_5$  represents a hydrogen- or a halogen atom in either the  $\alpha$ - or  $\beta$ -configuration;

$R_9$  represents a hydrogen atom or  $R_9$  represent a metal ion [e.g. The moiety  $SR_9$  represents a group of the formal  $[S]^- [M]^+$  wherein M is a metal such as Li, Na or K]; the method comprising;

A) reacting a steroidal carboxylic acid of formula (II) or a salt thereof



in which the substituents of formula (II) have the above defined meaning with a coupling agent alone or in conjunction with an coupling enhancer, followed by the reaction with a nucleophilic agent comprising a sulfur atom; and optionally

B) reacting the product from step A) with an acid.

8. (Currently Amended) The method of claim 1 ~~any of the preceding claims~~, wherein i) the coupling agent is added before the coupling enhancer, or the coupling enhancer is added before the coupling agent, and/or wherein ii) the steroidal carboxylic acid is added to a mixture of the coupling agent and the coupling enhancer, or wherein

a mixture of the coupling agent and the coupling enhancer is added to a steroidal

carboxylic acid, or wherein

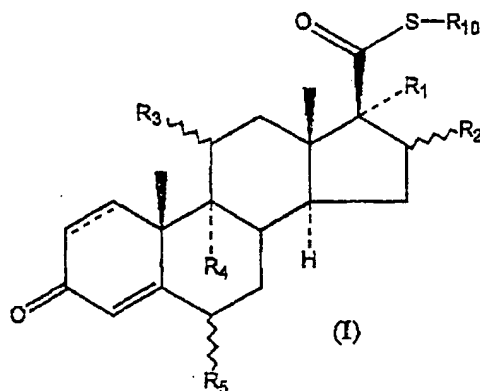
the steroidal carboxylic acid is added to a mixture of the coupling agent and the coupling enhancer in a polar aprotic solvent, preferably DMF or DMA, at elevated temperature.

9. (Currently Amended) A method for preparing a steroidal carbothioate (~~i.e. the ester of the steroidal carbothioic acid~~), or a salt thereof, the method comprising;

reacting a steroidal carbothioic acid or a salt thereof, ~~which is prepared as defined in any of the preceding claims,~~ with an electrophillic agent.

10. (Currently Amended) A method according to claim 9, in which the electrophillic agent is selected from the group consisting of: C<sub>1-8</sub> di- or trihaloalkanes, ~~preferably a trihalo- or a dihalomethane, such as chlorobromomethane or bromofluoromethane.~~

11. (Currently Amended) A method according to claim 9 ~~or 10~~ for preparing a steroidal carbothioate of formula (I)

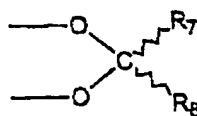


wherein R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub> and R<sub>5</sub> are defined as in claim 7;

R<sub>1</sub> represents a hydrogen atom, a hydroxy- or an alkoxy group in the  $\alpha$ -configuration, a group -O-C(=O)-R<sub>6</sub>, is an alkyl group or an optionally substituted 5-6 membered heterocyclic ring containing either oxygen, nitrogen or sulfur as ring hetero atom;

R<sub>2</sub> represents a hydrogen atom, a hydroxy group, an alkoxy group in the  $\beta$ -configuration,

an alkyl group which may be in either the  $\eta$ - or  $\beta$ -configuration, an alkylene group, wherein the alkylene group is bound to the steroid nucleus via a double bond, or  $R_1$  and  $R_2$  together represent



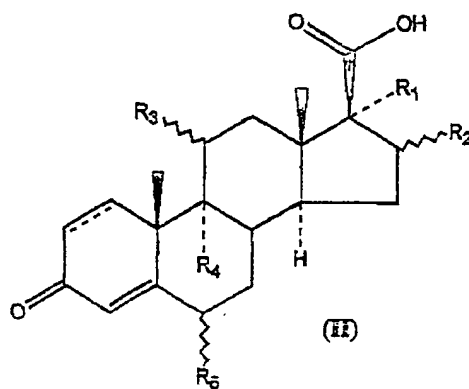
where  $R_7$  and  $R_8$  are the same or different and each represent a hydrogen atom or an alkyl group:

$R_3$  represent a hydrogen atom, hydroxy- or a protected hydroxy group in either a  $\alpha$ - or  $\beta$ -configuration or an oxo group:

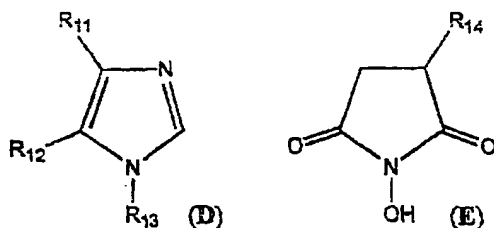
$R_4$  represents a hydrogen- or a halogen atom or  $R_3$  and  $R_4$  together represent a carbon-carbon bond or an epoxy group in the  $\beta$ -configuration: and

$R_5$  represents a hydrogen- or a halogen atom in either the  $\alpha$ - or  $\beta$ -configuration  
and  $R_{10}$  represents a  $C_{1-5}$  haloalkyl or an optionally substituted heterocyclic ring, the method comprising:

A) reacting a steroidal carboxylic acid of formula (II)



with a coupling agent and a coupling enhancer ~~[such as a compound of formula (D) or formula(E)]~~



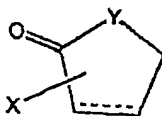
wherein  $R_{11}$  and  $R_{12}$  independently represent a hydrogen atom or a cyano group ( $C\equiv N$ );

$R_{12}$  represent a hydrogen atom or an alkyl group; and

$R_{14}$  represent a hydrogen atom or a moiety of a sulfonic acid ~~such as sodium sulfonate~~  
(e.g. The group  $-S(=O)(=O)-O-Na^+$ );

B) reacting the product from step A) with a nucleophilic agent comprising sulfur; and

C) reacting the product from step B) with an electrophilic agent ~~{such as a  $C_{1-6}$ - or trihaloalkane, preferably a trihalo- or a dihalomethane such as chlorofluoromethane or bromofluoromethane}~~ or a compound of the following formula;



wherein  $X=H, F, Cl, \text{ or } Br$  and;  $Y=CH_2, NH, O, \text{ or } S$ ; preferably  $X=Cl$  and  $Y=O$ .

12. (Original) The method of claim 11, wherein the coupling enhancer is selected from the group consisting of: NMI (N-methylimidazole); DCI (4,5-dicyanolimidazole); NHS (N-hydroxysuccinimide); and sulfo-NHS (N-hydroxysulfosuccinimide).

13. (Currently Amended) The method of ~~any of the claims 11-12~~, wherein step C) constitutes the *in situ* reaction of the product from step B) with bromofluoromethane to form a



compound of formula (I) wherein R<sub>10</sub> is a fluoromethyl group, such as fluticasone propionate.

14. (Currently Amended) The method according to claim 9 ~~any of the preceding claims~~, in which

at least two subsequent steps are performed *in situ*, ~~i.e. without any change or removal of solvents, or isolation of the individual intermediates; and/or~~

the method is conducted as a continuous method; ~~and/or~~

step A), B) and optionally step C) are conducted as a one-pot synthesis without solvent changes, ~~and/or~~ are performed at room or elevated temperature, or both; or

a combination of one or more of the foregoing.

15. (Currently Amended) The method of ~~any of the claims 9-14~~, wherein an androstane 17 $\beta$ -carboxylic acid is converted to an androstane 17 $\beta$ -carbothioate.

16. (Currently Amended) The method of claim 9 ~~any of the preceding claims~~, wherein step B) provides ~~an alkali metal salt of the thioic acid, such as a compound of formula (IV), in which the moiety -S-R<sub>5</sub> represent a group of the formula [-S]<sup>-</sup>[M]<sup>+</sup> wherein M is a metal such as Li, Na or K-e.g. S<sup>-</sup>Na<sup>+</sup>, and the other substituents have the same meaning as defined in claim 7.~~

wherein the symbol  $\equiv$  in the 1,2-position represent a single or a carbon-carbon double bond;

R<sub>1</sub> represents a hydrogen atom, a hydroxy- or an alkoxy group in the  $\alpha$ -configuration, a group -O-C(=O)-R<sub>6</sub> is an alkyl group or an optionally substituted 5-6 membered heterocyclic ring containing either oxygen, nitrogen or sulfur as ring hetero atom;

R<sub>2</sub> represents a hydrogen atom, a hydroxy group, an alkoxy group in the  $n$ -configuration, an alkyl group which may be in either the  $\eta$ - or  $\beta$ -configuration, an alkylene group, wherein the alkylene group is bound to the steroid nucleus via a double bond, or R<sub>1</sub> and R<sub>2</sub> together represent

where  $R_7$  and  $R_8$  are the same or different and each represent a hydrogen atom or an alkyl group;

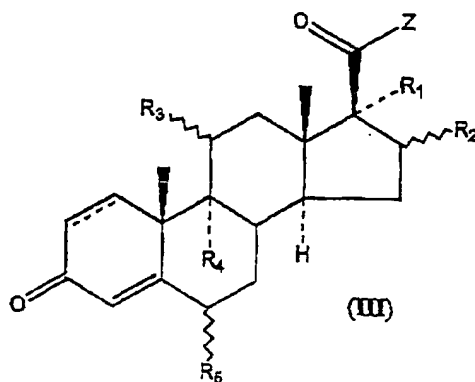
$R_3$  represent a hydrogen atom, hydroxy-or a protected hydroxy group in either a  $\alpha$ - or  $\beta$ -configuration or an oxo group;

$R_4$  represents a hydrogen- or a halogen atom or  $R_3$  and  $R_4$  together represent a carbon-carbon bond or an epoxy group in the  $\beta$ -configuration; and

$R_5$  represents a hydrogen- or a halogen atom in either the  $\alpha$ - or  $\beta$ -configuration;

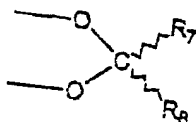
$R_9$  represents a hydrogen atom or  $R_9$  represent a metal ion.

17. (Currently Amended) A compound of the formula (III) and salts and solvates thereof



Wherein R<sub>1</sub> represents a hydrogen atom, a hydroxy- or an alkoxy group in the α-configuration, a group -O-C(=O)-R<sub>6</sub> is an alkyl group or an optionally substituted 5-6 membered heterocyclic ring containing either oxygen, nitrogen or sulfur as ring hetero atom;

R<sub>2</sub> represents a hydrogen atom, a hydroxy group, an alkoxy group in the η-configuration, an alkyl group which may be in either the η- or β-configuration, an alkylene group, wherein the alkylene group is bound to the steroid nucleus via a double bond, or R<sub>1</sub> and R<sub>2</sub> together represent



where R<sub>7</sub> and R<sub>8</sub> are the same or different and each represent a hydrogen atom or an alkyl group;

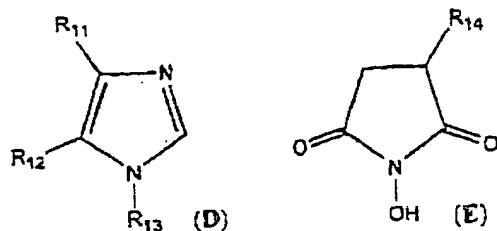
R<sub>3</sub> represent a hydrogen atom, hydroxy- or a protected hydroxy group in either a α- or β-configuration or an oxo group;

R<sub>4</sub> represents a hydrogen- or a halogen atom or R<sub>3</sub> and R<sub>4</sub> together represent a carbon-carbon bond or an epoxy group in the β-configuration; and

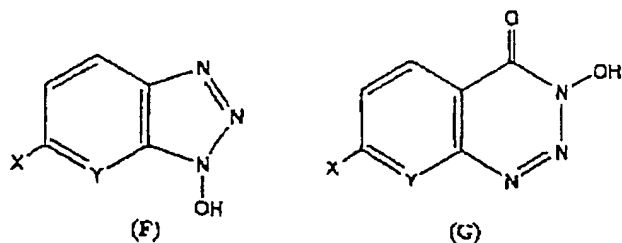
R<sub>5</sub> represents a hydrogen- or a halogen atom in either the α- or β-configuration; and

Z represent the structural moiety resulting from the reaction between the steroidal carboxylic acid of formula (II) and a coupling agent (preferably EDC), followed by a coupling

enhancer selected from the group consisting of the compounds of formulas (D); (E); (F); and (G):



wherein  $R_{11}$  and  $R_{12}$  independently represent a hydrogen atom or a cyano group;  $R_{13}$  represent a hydrogen atom or a methyl group; and  $R_{14}$  represent a hydrogen atom or a moiety of a sulfonic acid, such as sodium sulfonate [i.e. The group  $S(-O)(-O)O^-Na^+$ ],



$X = H, F, Cl, Br$  and  $Y = CH, N, O, S$

with the proviso that:

when the coupling enhancer is a compound of formula (F), X can not represent H when Y represents CH:

when the coupling enhancer is a compound of formula (D),  $R_{11}$  and  $R_{12}$  can not both represent H when  $R_1$  in formula III represents DH; and

when the coupling enhancer is a compound of formula (E),  $R_{14}$  can not represent H when  $R_1$  in formula III represents H;

and with the further proviso that

succinimidyl-9 $\alpha$ -fluoro-11 $\beta$ , 17 $\alpha$ -dihydroxy-16 $\alpha$ -methyl-3-oxoandrosta-1,4-diene-17 $\beta$ -carboxylate;

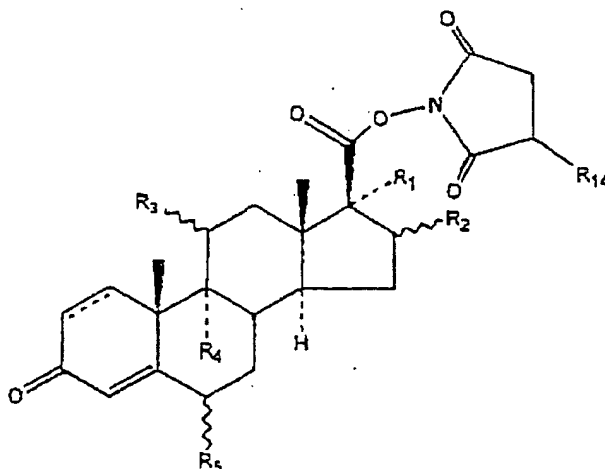
17 $\alpha$ -hydroxy-4-androsten-3-one-17 $\beta$ -carboxylic acid N-hydroxysuccinimide ester;

N-hydroxysuccinimidyl-9-fluoro-16 $\alpha$ -methyl-11 $\beta$ , 17-dihydroxy-3-oxo-1,4-androstadiene-17 $\beta$ -carboxyester;

N-hydroxysuccinimide ester of dexamethasone-17 $\beta$ -carboxylic acid; and 1-[(9-fluoro-11 $\beta$ -hydroxy-16 $\beta$ -methyl-3-oxo-17 $\alpha$ -propionylaxyandrosta-1,4-dien-17 $\beta$ -yl)carbonyl]imidazol are disclaimed.

18. (Currently Amended) The compound of claim 17, wherein at least one of R<sub>11</sub> and R<sub>12</sub> is a cyano group (C=N), and/or R<sub>13</sub> is a hydrogen atom, and/or formula (D) is NMI (N-methylimidazole) or DCI (4,5-dicyano-imidazole), and/or formula (E) is NHS (N-hydroxysuccinimide) or sulfo-NHS (N-hydroxysulfosuccinimide), or a combination comprising one or more of the foregoing.

19. (Currently Amended) The compound of claim 17, having the formula:



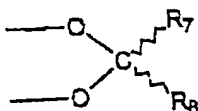
~~in which the substituents have the same meaning as defined in claim 17, and salts and solvates thereof, with the proviso that R<sub>14</sub> can not represent H when R<sub>1</sub> represents H.~~

20.(Currently Amended) A compound of the formula (VI) and salts and solvates thereof

wherein  $R_1$ ,  $R_2$ ,  $R_3$ ,  $R_4$  and  $R_5$  are defined as in claim 7, and  $R_a$  and  $R_b$  are defined as in claim 1;

$R_1$  represents a hydrogen atom, a hydroxy- or an alkoxy group (such as an optionally substituted  $C_{1-6}$  alkoxy) in the  $\alpha$ -configuration, a group  $-O-C(=O)-R_6$  is an alkyl group (such as optionally substituted  $C_{1-6}$  alkyl) or an optionally substituted 5-6 membered heterocyclic ring containing either oxygen, nitrogen or sulfur as ring hetero atom (such as a furanyl, pyrrolyl or a thiophenyl group);

$R_2$  represents a hydrogen atom, a hydroxy group, an alkoxy group (such as an optionally substituted  $C_{1-6}$  alkoxy) in the  $n$ -configuration, an alkyl group (such as an optionally substituted  $C_{1-6}$  alkyl) which may be in either the  $\eta$ - or  $\beta$ -configuration, an alkylene group (such as an optionally substituted  $C_{1-6}$  alkylene having the two free valencies on the same carbon atom preferably methylene), wherein [the alkylene group is bound to the steroid nucleus via a double bond,] or  $R_1$  and  $R_2$  together represent



where  $R_7$  and  $R_8$  are the same or different and each represent a hydrogen atom or an alkyl group (such as an optionally substituted  $C_{1-6}$  alkyl);

$R_3$  represent a hydrogen atom, hydroxy- or a protected hydroxy group in either a  $\alpha$ - or  $\beta$ -configuration or an oxo group (in which case the bond between  $R_3$  and the steroid nucleus is a double bond);

$R_4$  represents a hydrogen- or a halogen atom or  $R_3$  and  $R_4$  together represent a carbon-carbon bond or an epoxy group in the  $\beta$ -configuration; and

$R_5$  represents a hydrogen- or a halogen atom in either the  $\alpha$ - or  $\beta$ -configuration,

wherein  $R_a$  and  $R_b$  are the same or different, and each represent an aliphatic, heteroaliphatic, carbocyclic or a heterocyclic group;

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with the proviso that 1-(3-dimethylamino-propyl)-3-ethyl-carbodiimide-6 $\alpha$ , 9 $\alpha$ -difluoro-11 $\beta$ -hydroxy-16 $\alpha$ , 17 $\alpha$ -isopropylidenedioxy-3-oxo-androsta-1,4-diene-17 $\beta$ -carboxylate is disclaimed.

21-23. (Cancelled).